PTO/SB/08A (08-03) Approved for use through 07/31/2006. OMB 0651-0031

U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE
Under the Paperwork Reduction Act of 1995 for persons are required to respond to a collection of information unless it contains a valid OMB control number.

Substitute for form 1449A/PTO

TRADEILA PR

INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(use as many sheets as necessary)

Sheet 1 of 6

	Complete if Known
Application Number	10/602,142
Filing Date	June 20, 2003
First Named Inventor	Sommadossi et al.
Group Art Unit	Unassigned
Examiner Name	Unassigned
Attorney Docket Number	06171.105076 IDX 1007 CON2

						3425610_1			
U.S. PATENT DOCUMENTS									
Examiner Initials *	Cite No. 1	U.S. Patent Docu Number	ment Kind Code if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pgs, Clmns, Lns, Where Relevant Passages/Relevant Figs Appear			
40	AA	3,480,613	Α	Walton et al.	11-25-1969				
1	AB	5,977,061	A	De Clercq	11-02-1999				
	AC	6,340,690	B1	Bachand et al.	01-22-2002				
	AD	6,348,587	Bl	Schinazi et al.	02-2002				
	AE	6,395,716	B1	Gosselin et al. (Novirio / Idenix)	05-28-2002				
	AF	6,444,652	Bl	Gosselin et al. (Novirio / Idenix)	09-03-2002				
	AG	6,573,248	Bi	Ramasamy et al.	06-03-2003				
	AH	2002/0019363	Al	Ismaili et al.	02-2002				
	Αl	2002/0055483	Al	Watanabe et al.	05-09-2002				
	AJ	2002/0147160	Al	Bhat et al.	10-10-2002				
	AK	2003/008841	Al	Devos et al.	01-09-2003				
	AL	2003/028013	Al	Wang et al.	02-06-2003	•			
	AM	2003/0050229	Al	Sommadossi et al.	03-13-2003				
	AN	2003/0060400	Al	LaColla et al.	03-27-2003				
	AO	2003/0083307	A1	Devos et al.	05-01-2003				
A 410	AP	2003/0087873	Al	Stuyver et al.	05-08-2003				

					EIGN PATENT DOCUMENTS			
r	G:.	Fore	ign Patent Docun	nent		Date of	Pages, Columns,	
Examiner Cite Initials * No.		Office ³ Number Kind Code ² (if known)			Name of Patentee or Applicant of Cited Document	Publication of Cited Document MM-DD- YYYY	Lines, Where Relevant Passages or Relevant Figures Appear	T⁵
	AQ	FR	1,521,076	A	Merck & Co. Inc.	04-12-1968		
	AR	FR_	1,581,628	A	Merck & Co. Inc.	09-19-1969		
	AS	FK	2,662,165	A	Univ. Paris Curie	11-22-1991		
10	AT	GB .	1,163,103	Α	Merck & Co. Inc.	09-04-1969		
40	AU	GB	1,209,654	Α	Merck & Co. Inc.	10-21-1970		
	AV-	JP-	-63-215694	A	Yamasa Shoyu Co. Ltd.	09-08-1988		
,	AW	JP.	06-228186	-A	Yamasa Shoyu Co. Ltd.	08-16-1994		
Kt.O	AX	WO	98/16184	A2	ICN Pharmaceuticals.	04-23-1998		
χ̈́χ	AY	WO	99/43691	Al	Emory U.; U.Ga.R.F.	02-09-1999		\Box
100	AZ	WO	00/09531	A2	Novirio Pharm. (Idenix)	02-24-2000		
NU	AAA	WO	01/32153	A2	Biochem Pharma	05-10-2001		

Character W. C.	onsidered 3/6/05
---	------------------

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

Burden Hour Statement: This form is estimated to take 2.0 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, U. S. Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Washington, DC 20231.

¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

Approved for use through 07/31/2006. OMB 0651-0031

U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

Complete if Known Substitute for form 1449A/PTO **Application Number** 10/602,142 Filing Date INFORMATION DISCLOSURE June 20, 2003 First Named Inventor STATEMENT BY APPLICANT Sommadossi et al. Group Art Unit Unassigned Examiner Name (use as many sheets as necessary) Unassigned Attorney Docket Number 6 Sheet 06171.105076 IDX 1007 CON2

							342561	IU I
,,	·	-	 		EIGN PATENT DOCUMENTS	-	,	
Examiner Initials *	Cite No. 1		ign Patent Docur Number Kind (if k		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD- YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁶
HO	BA	wo	01/60315	A2	Biochem Pharma	08-23-2001		Г
	BB	wo	01/68663	Al	ICN Pharmaceuticals	09-20-2001		
	BC	WO	01/79246	A2	Pharmasset	10-25-2001		
	BD	wo	01/90121	A2	Novirio Pharm. (Idenix)	11-29-2001		
	BE	WO	01/91737	A2	Novirio Pharm. (Idenix)	06-12-2001		
	BF	wo	01/92282	A2	Novirio Pharm. (Idenix)	06-12-2001		
	BG	WO	01/96353	A2	Novirio Pharm. (Idenix)	12-20-2001		1
	ВН	WO	02/03997	Al	ICN Pharmaceuticals	01-17-2002		
	BI	WO	02/18404	A2	F. Hoffmann-La Roche	03-07-2002		一
	BJ	WO	02/32920	A2	Pharmasset	04-25-2002		
	BK	wo	02/48165	A2	Pharmasset	06-20-2002		┼─
	BL	wo	02/057287	A2	Merck & Co. Inc.	07-25-2002		\vdash
	BM	wo	02/057425	A2	Merck & Co. Inc.	07-25-2002		Т
	BN	wo	02/070533	A2	Pharmasset	09-12-2002		\vdash
	BO	wo	02/094289	Al	F. Hoffmann-La Roche	11-28-2002		
	BP	wo	02/100415	A2	F. Hoffmann-La Roche	12-19-2002		
	BQ	wo	03/026589	A2	Idenix; CNRS; U. Montp.	04-03-2003		
	BR	wo	03/026675	A1	Idenix; CNRS; U. Montp.	04-03-2003		
	BS	wo	03/051899	Al	Ribapharm	06-26-2003		-
	BT	wo	03/061385	Al	Ribapharm	07-31-2003		\vdash
	BU	wo	03/061576	A2	Ribapharm	07-31-2003		\vdash
	BV	wo	03/062255	A2	Ribapharm	07-31-2003		\vdash
	BW	wo	03/062256	A1	Ribapharm	07-31-2003		\vdash
	ВХ	wo	03/062257	Al	Ribapharm	07-31-2003		
	BY	WO	03/063771	A2	Pharmasset	08-07-2003		
	BZ	wo	03/068162	A2	Pharmasset	08-21-2003		
	BAA	wo	03/072757	A2	Biota Inc.	09-04-2003		_
	BAB	wo	03/093290	A2	Genelabs Technologies	11-13-2003		-
1 11	BAC	wo	04/002422	A2	Idenix; Univ.D.S.Cagliari	01-08-2004		
OFF	BAD	wo	04/002999	A2	Idenix; Univ.D.S.Cagliari	01-08-2004		_

Examiner Date		
Signature Consideration Consid	red 3/2	1/05

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant,

¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ³ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. Applicant is to place a check mark here if English language Translation is attached.

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number

					The state of the s	
Substitut	e for form 1449A/PTO			Complete if Known		
Substitut	e to tom 1449AFTO			Application Number	10/602,142	
INF	ORMATION D	ISCL	OSURE	Filing Date	June 20, 2003	
STA	TEMENT BY	APPL	CANT	First Named Inventor	Sommadossi et al.	
l				Group Art Unit	Unassigned	
	(use as many sheets	as necessai	אי	Examiner Name	Unassigned	
Sheet	3	of	6	Attorney Docket Number	06171.105076 IDX 1007 CON2	

3425610 1 OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, Examiner Cite No. 1 Initials * journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. ALTMANN et al, "The synthesis of 1'-methyl carbocyclic thymidine and its effect on nucleic acid duplex stability," Synlett, Thieme Verlag, Stuttgart, De, 10:853-855 (1994). CB BAGINSKI, S. G, et al., "Mechanism of action of a pestivirus antiviral compound," PNAS USA. 97(14):7981-7986 (2000). BEIGELMAN, L.N., et al, "Epimerization during the acetolysis of 3-O-acetyl-5-O-benzoyl-1,2-Oisopropylidene-3-C-methyl-α,D-ribofuranose. Synthesis of 3'-C-methylnucleosides with the β-Dribo- and α-D-arabino configurations," Carbohydrate Research, 181:77-88 (1988). CD BEIGELMAN, L.N., et al, "A general method for synthesis of 3'-C-alkylnucleosides," Nucleic Acids Symp. Ser., 9:115-118 (1981). CE BERENGUER, M., et al, "Hepatitis B and C viruses: Molecular identification and targeted antiviral therapies," Proceedings of the Association of American Physicians, 110(2), 98-112 (1998). CARROLL, S.S., et al., "Inhibition of hepatitis C virus RNA replication by 2'-modified nucleoside CF analogs," The Journal of Biological Chemistry, 278(14):11979-11984 (2003). CG CZERNECKI, S., et al, "Synthesis of various 3'-branched 2',3'-unsaturated pyrimidine nucleosides as potential anti-HIV agents," J. Org. Chem., 57:7325-7328 (1992). CH De FRANCESCO, R., et al., "Approaching a new era for hepatitis C virus therapy: Inhibitors of the NS3-4A serine protease and the NS5B RNA-dependent RNA polymerase," Antiviral Research, 58:1-16 (2003). CI FAIVRE-BUET, V., et al, "Synthesis of 1'-deoxypsicofuranosyl-deoxynucleosides as potential anti-HIV agents," Nucleosides & Nucleotides, 11(7):1411-1424 (1992). CJ FARKAS, J., et al., "Nucleic acid components and their analogues. XCIV. Synthesis of 6-amino-9-(1deoxy-β-D-psicofuranosyl)purine", Collect. Czech. Chem. Commun. 32:2663-2667 (1967). CK FARKAS, J., et al., "Nucleic acid components and their analogues. LXXIX. Synthesis of methyl 1deoxy-D-psicofuranosides substituted at C(1) with halo atoms or a mercapto group," Collect. Czech. Chem. Commun., 31:1535-1543 (1996). FEDOROV, I.I., et al, "3'-C-Branched 2'-deoxy-5-methyluridines: Synthesis, enzyme inhibition, and CL antiviral properties," J. Med. Chem., 35(24):4567-4575 (1992). FRANCHETTI, P., et al., "2'-C-Methyl analogues of selective adenosine receptor agonists: synthesis CM and binding studies," J. Med. Chem., 41(10):1708-1715 (1998). GROUILLER, A., et al., "Novel p-toluenesulfonylation and thionocarbonylation of unprotected thymine nucleosides," Synlett, 1993, 221-222 (March 1993). CO HARAGUCHI, K., et al., "Preparation and reactions of 2'- and 3'- vinyl bromides of uracil nucleosides: Versatile synthons for anti-HIV agents," Tetrahedron Letters, 32(28):3391-3394 (1991).

	The same of the sa		
Examiner Signature	Hadleds	Date Considered	3/6/05
The second secon			

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ³ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

Approved for use through 07/31/2006. OMB 0651-0031
U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE
o a collection of information unless it contains a valid OMB control number.

Substitute for	form 1449A/PTO				Complete if Known
Substitute for	101111 147701 10			Application Number	10/602,142
INFOR	RMATION	DISCLO	SURE	Filing Date	June 20, 2003
STATEMENT BY APPLICANT			CANT	First Named Inventor	Sommadossi et al.
				Group Art Unit	Unassigned
(use as many sheets as necessary))	Examiner Name	Unassigned
Sheet	4	of	6	Attorney Docket Number	06171.105076 IDX 1007 CON2

		34256	510_1
	,	OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
Examiner	Cite	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine,	
Initials *	No. 1	journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T°
VA	DA	HARAGUCHI, K., et al., "Stereoselective synthesis of 1'-C-branched uracil nucleosides from	
No		uridine," Nucleosides & Nucleotides, 14(3-5):417-420 (1995).	
1	DB	HARRY-O'KURU, R.E., et al., "A short, flexible route toward 2'-C-branched ribonucleosides",	
		J.Org. Chem., 62:1754-1759 (1997). (Scheme 11).	
	DC	HARRY-O'KURU, R.E., et al., "2'-C-Alkylribonucleosides: Design, synthesis, and conformation,"	
		Nucleosides & Nucleotides, 16(7-9):1457-1460 (1997). ["Rogers" in #2; correct name in #7]	
	DD	HATTORI, H., et al, "Nucleosides and nucleotides. 175. Structural requirements of the sugar moiety	
ł		for the antitumor activities of new nucleoside antimetabolites, 1-(3-C-ethynyl-b-D-ribo-	
		pentofuranosyl)cytosine and -uracil," J. Med. Chem., 41:2892-2902 (1998).	
	DE	HREBABECKY, H., et al., "Nucleic acid components and their analogues. CXLIX. Synthesis of	
		pyrimidine nucleosides derived from 1-deoxy-D-psicose," Collect. Czech. Chem. Commun., 37:2059-	
		2065 (1972).	
	DF	HREBABECKY, H., et al. "Synthesis of 7- and 9β-D-psicofuranosylguanine and their 1'-deoxy	-
derivatives," Collect. Czech. Chem. Commun., 39:2115-2123 (1974).			
	DG IINO, T., et al., "Nucleosides and nucleotides. 139. Stereoselective synthesis of (2'S)-2'-C-alky		
	1	deoxyuridines," Nucleosides and Nucleotides, 15(1-3):169-181 (1996).	
	DH	ITOH, Y., et al, "Divergent and stereocontrolled approach to the synthesis of uracil nucleosides	
		branched at the anomeric position," J. Org. Chem., 60(3):656-662 (1995).	
	DI	JOHNSON, C.R., et al, "3'-C-Trifluoromethyl ribonucleosides," Nucleosides & Nucleotides,	
i	}	14(1&2):185-194 (1995).	
	DJ	KAWANA, M., et al., "The deoxygenation of tosylated adenosine derivatives with Grignard	_
	ĺ	reagents," Nucleic Acids Symp. Ser., 17:37-40 (1986).	
	DK	LAVAIRE, S., et al., "3'-Deoxy-3'-C-trifluoromethyl nucleosides: Synthesis and antiviral	
		evaluation," Nucleosides & Nucleotides, 17(12):2267-2280 (1998).	
	DL	LEYSSEN, P. et al., "Perspectives for the treatment of infections with Flaviviridae," Clinical	
		Microbiology Reviews (Washington, D.C.), 13(1):67-82 (January 2000).	
	DM	MARTIN, X., et al., "Intramolecular hydrogen bonding in primary hydroxyl of thymine 1-(1-deoxy-β-	
		D-psicofuranosyl) nucleoside," Tetrahedron, 50(22):6689-6694 (1994).	
	DN	MATSUDA, A., et al., "Radical deoxygenation of tert-alcohols in 2'-branched-chain sugar pyrimidine	
	J.,	nucleosides: Synthesis and antileukemic activity of 2'-deoxy-2'(S)-methylcytidine," Chem. Pharm.	
		Bull., 35(9):3967-3970 (1987).	
	DO	MATSUDA, A., et al., "Alkyl addition reaction of pyrimidine 2'-ketonucleosides: Synthesis of 2'-	
Mh		branched-chain sugar pyrimidine nucleosides (Nucleosides and Nucleotides. LXXXI)," Chem.	
.Alo		Pharm. Bull., 36(3):945-953 (1988).	
		1 marm. Dam, 30(3).773-733 (1700).	

		The state of the s
Examiner Signature	Date Considered	3/6/05

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the rign of the Emperor must precede the serial number of the patent document. ³ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

	. C			Complete if Known		
Substitute	e for form 1449A/PTO			Application Number	10/602,142	
INF	ORMATION D	ISCL	OSURE	Filing Date	June 20, 2003	
STATEMENT BY APPLICANT				First Named Inventor	Sommadossi et al.	
				Group Art Unit	Unassigned	
	(use as many sheets	as necessar)	Examiner Name	Unassigned	
Sheet	5	of	6	Attorney Docket Number	06171.105076 IDX 1007 CON2	

3425610 1 OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, Examiner Cite Initials * No. I journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. MATSUDA, A., et al., "Nucleosides and Nucleotides. 94. Radical deoxygenation of tert-alcohols in 1-(2-C-alkylpentofuranosyl)pyrimidines: Synthesis of (2'S)-2'-deoxy-2'-C-methylcytidine, an antileukemic nucleoside, "J. Med. Chem., 34:234-239 (1991). MATSUDA, A., et al., "Nucleosides and Nucleotides, 104. Radical and palladium-catalyzed **EB** deoxygenation of the allylic alcohol systems in the sugar moiety of pyrimidine nucleosides," Nucleosides & Nucleotides, 11(2/4):197-226 (1992). MIKHAILOV, S.N., et al., "Synthesis and properties of 3'C-methylnucleosides and their phosphoric EC esters," Carbohydrate Research, 124:75-96 (1983). MIKHAILOV, S.N., et al., "Substrate properties of C'-methylnucleoside and C'-methyl-2'-ED deoxynucleoside 5'-triphosphates in RNA and DNA synthesis reactions catalysed by RNA and DNA polymerases," Nucleosides & Nucleotides, 10(1-3):339-343 (1991). MIKHAILOV, S.N., et al, "Hydrolysis of 2'- and 3'-C-methyluridine 2'c3'-cyclic monophosphates EE and interconversion and dephosphorylation of the resulting 2'- and 3'-monophosphates: Comparison with the reactions of uridine monophosphates," J. Org. Chem., 57 (15):4122-4126 (1992). EF NUTT, R.F., et al., "Branched-chain sugar nucleosides. III. 3'-C-methyladenine", J.Org. Chem., 33:1789-1795 (1968). OIVANEN, M., et al, "Additional evidence for the exceptional mechanism of the acid-catalyzed hydrolysis of 4-oxopyrimidine nucleosides: Hydrolysis of 1-(1-alkoxyalkyl)uracils, seconucleosides, 3'-C-alkyl nucleosides and nucleoside 3',5'-cyclic monophosphates," J. Chem. Soc. Perkin Trans. 2, 1994:309-314 (1994). ONG, S.P., et al, "Synthesis of 3'-C-methyladenosine and 3'-C-methyluridine diphosphates and their interaction with the ribonucleoside diphosphate reductase from Corynebacterium nephridii," Biochemistry, 31(45):11210-11215 (1992). EI Oral Session V, Hepatitis C Virus, Flaviviridae; 16th International Conference on Antiviral Research (April 27, 2003, Savannah, Ga.) p A75-77. EJ PAN-ZHOU, X-R, et al., "Differential effects of antiretroviral nucleoside analogs on mitochondrial function in HepG2 cells," Antimicrob. Agents Chemother., 44:496-503 (2000). EK ROSENTHAL, A., et al., "Branched-chain sugar nucleosides. Synthesis of 3'-C-ethyl (and 3'-Cbutyl)uridine Carbohydrate Research, 79:235-242 (1980). EL SAMANO, V., et al., "Synthesis and radical-induced ring-opening reactions of 2'-deoxyadenosine-2'spirocyclopropane and its uridine analogue. Mechanistic probe for ribonucleotide reductases," J. Am. Chem. Soc., 114:4007-4008 (1992).

Examiner Signature	Date Considered 36/05
	Mojos

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ³ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

Substitute for form 1449A/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)				Complete if Known		
				Application Number	10/602,142	
			OSURE	Filing Date	June 20, 2003	
			ICANT	First Named Inventor	Sommadossi et al.	
				Group Art Unit	Unassigned	
			אמ	Examiner Name	Unassigned	
Sheet	6	of	6	Attorney Docket Number	06171.105076 IDX 1007 CON2	

3425610 1 OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, Examiner Cite No. 1 Initials • journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. FA SAMANO, V., et al., "Nucleic acid related compounds. 77. 2',3'-Didehydro-2',3'-dideoxy-2'(and 3')-methylnucleosides via [3,3]-sigmatropic rearrangements of 2'(and 3')-methylene-3'(and 2')-Othiocarbonyl derivatives and radical reduction of a 2'-chloro-3'-methylene analogue," Can. J. Chem. 71:186-191 (1993). SCHMIT, C., et al, "The effects of 2'- and 3'-alkyl substituents on oligonucleotide hybridization and stability," Biorganic & Medicinal Chemistry Letters, 4(16):1969-1974 (1994). ["Altmann"] FC SERAFINOWSKI, P.J., et al., "New method for the preparation of some 2'- and 3'-trifluoromethyl-2',3'-dideoxyuridine derivatives," Tetrahedron (Elsevier Science Publishers), 56(2):333-339 (1999). FD SHARMA, P.K., et al., "Synthesis of 3'-trifluoromethyl nucleosides as potential antiviral agents." Nucleosides, Nucleotides and Nucleic Acids, 19(4):757-774 (2000). FE SOMMADOSSI J-P, et al., "Comparison of cytotoxicity of the (-)- and (+)-enantiomer of 2',3'dideoxy-3'-thiacytidine in normal human bone marrow progenitor cells" Biochemical Pharmacology, 44:1921-1925 (1992). FF SOMMADOSSI J-P, et al., "Toxicity of 3'-azido-3'-deoxythymidine and 9-(1,3-dihydroxy-2propoxymethyl)guanine for normal human hematopoietic progenitor cells in vitro" Antimicrobial Agents and Chemotherapy, 31:452-454 (1987). TRITSCH, D., et al., "3'-β-ethynyl and 2'-deoxy-3'-β-ethynyl adenosines: First 3'-β-branched adenosines substrates of adenosine deaminase," Bioorganic & Medicinal Chemistry Letters, 10:139-TUNITSKAYA, V.L., et al., "Substrate properties of C'-methyl UTP derivatives in T7 RNA polymerase reactions. Evidence for N-type NTP conformation," FEBS Letters, 400:263-266 (1997). FI USUI, H., et al., "Synthesis of 2'-deoxy-8,2'-ethanoadenosine and 3'-deoxy-8,3'-ethanoadenosine (Nucleosides and Nucleotides. LXIV)," Chem. Pharm. Bull., 34(1):15-23 (1986). WALCZAK, K., et al., "Synthesis of 1-(3-alkyl-2,3-dideoxy-D-pentofuranosyl)uracils with potential FJ anti-HIV activity," Acta Chemica Scand., 45:930-934 (1991). FK WALTON, E., et al., "Branched-chain sugar nucleosides. V. Synthesis and antiviral properties of several branched-chain sugar nucleotides," J. Med. Chem., 12:306-309 (1969). WOLFE, M.S., et al., "A concise synthesis of 2'-C-methylribonucleosides," Tetrahedron Letters, FL 36(42):7611-7614 (1995). WU, J.-C., et al., "A new stereospecific synthesis of [3.1.0] bicyclic cyclopropano analog of 2',3'-**FM** dideoxyuridine, Tetrahedron, 46(7):2587-2592 (1990).

Examiner	2// 2//	Date/	1
Signature	Horl (S	Considered 2//	65

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.